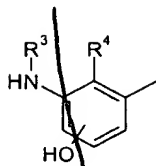


*Sub B1*



R<sup>1</sup> is

wherein:

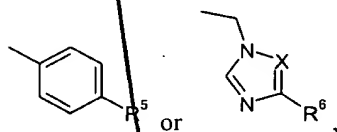
R<sup>3</sup> is a benzyl group optionally substituted by a methoxy group,

R<sup>4</sup> is a hydrogen atom, or

R<sup>3</sup> and R<sup>4</sup> together are a -CO-CH<sub>2</sub>-O- bridge, the carbonyl group of the bridge being bound to the nitrogen; and

*A1 cont*

R<sup>2</sup> is



wherein:

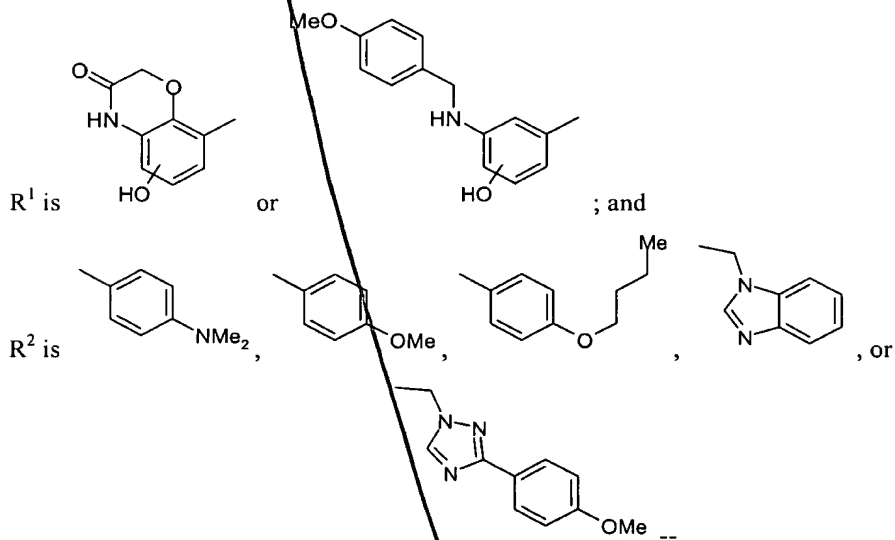
R<sup>5</sup> is a dimethylamino, methoxy, or butoxy group,

X is a nitrogen or a carbon atom, and

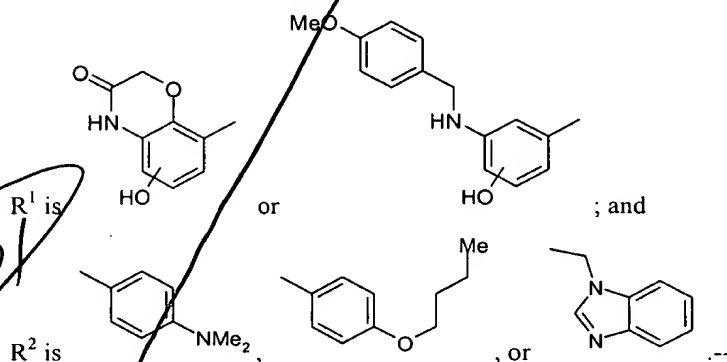
R<sup>6</sup> is a methoxyphenyl group, if X is nitrogen, or is an anellated phenyl ring also linked to X, if X is carbon,

or acid addition salt thereof.--

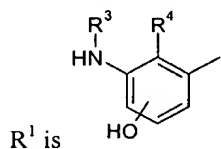
--2. (Amended) The compound of formula 1 according to claim 1, wherein:



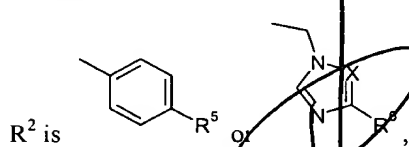
--3. (Amended) The compound of formula 1 according to claim 1, wherein:



--4. (Amended) The compound of formula 1 according to claim 1, wherein:



wherein  $R^3$  and  $R^4$  together are a  $-\text{CO}-\text{CH}_2-\text{O}-$  bridge, the carbonyl group of the bridge being bound to the nitrogen, and



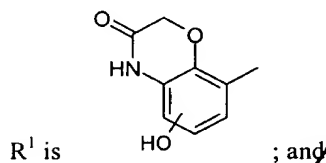
wherein:

$R^5$  is a dimethylamino, methoxy, or butoxy group,

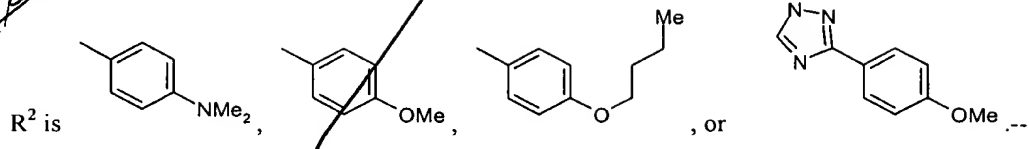
X is a nitrogen or a carbon atom, and

$R^6$  is a methoxyphenyl group, if X is nitrogen, or an anellated phenyl ring also linked to X, if X is carbon.--

--5. (Amended) The compound of formula 1 according to claim 1, wherein:

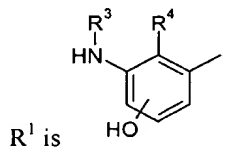


*Sub B2*



*A 2*  
*cont*

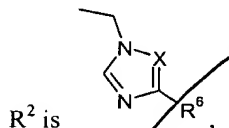
--6. (Amended) The compound of formula 1 according to claim 1, wherein:



wherein:

R<sup>3</sup> is a benzyl group optionally substituted by methoxy, and

R<sup>4</sup> is a hydrogen atom; and



wherein:

X is a nitrogen or a carbon atom,

R<sup>6</sup> is a methoxyphenyl group, if X is nitrogen, or an anellated phenyl ring also linked to

X, if X is carbon.

*A 2*

--8. (Amended) 1-[3-(4-methoxybenzylamino)-4-hydroxyphenyl]-2-[4-(1-benzimidazolyl)-2-methyl-2-butylamino]ethanol, or an acid addition salt thereof.--

--9. (Amended) 1-[2H-5-hydroxy-3-oxo-4H-1,4-benzoxazin-8-yl]-2-[3-(4-N,N-dimethylaminophenyl)-2-methyl-2-propylamino]ethanol, or an acid addition salt thereof.--

--10. (Amended) 1-[2H-5-hydroxy-3-oxo-4H-1,4-benzoxazin-8-yl]-2-[3-(4-n-butyloxyphenyl)-2-methyl-2-propylamino]ethanol, or an acid addition salt thereof.--

--11. (Amended) The compound according to one of claims 1 to 6 or 8 to 10, wherein the acid addition salt thereof is formed with a pharmacologically acceptable acid.--

*Sub B3*

A2  
Ced  
Sub 3  
--12. (Amended) A method of treating bronchial asthma, the inflammatory component in COPD, premature onset of labor in midwifery (tocolysis), atrio-ventricular block, bradycardiac hearth rhythm disorders, circulatory shock, or itching and inflammation of the skin in a host in need of such treatment, the method comprising administering to the host the compound according to one of claims 1 to 6 or 8 to 10.--

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--13. (Amended) A pharmaceutical preparation comprising a compound according to one of claims 1 to 6 or 8 to 10 and a conventional excipient or carrier.--

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A3  
--16. (New) The compound according to claim 7, wherein the acid addition salt thereof is formed with a pharmacologically acceptable acid.--

--17. (New) A pharmaceutical preparation comprising a compound according claim 7 and a conventional excipient or carrier.--

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Sub 4